

UNITED STATES PATENT AND TRADEMARK OFFICE

ENTTED STATES DEPARTMENT OF COMMERCE Enited States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Vignita 22313-1450 www.usplo.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
09/881,012	06/13/2001	Edward I. Ginns	015280-248120US	8624		
75	7590 07:06/2004			EXAMINER		
VENABLE, BAETJER, HOWARD & CIVILETTI, LLP			GOLDBERG, JEANINE ANNE			
1201 NEW YORK AVE, N.W. SUITE 1000		ART UNIT	PAPER NUMBER			
WASHINGTON, DC 20005			1634			

DATE MAILED: 07/06/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
Advisory Action	09/881,012	GINNS ET AL.				
Advisory Addon	Examiner	Art Unit				
	Jeanine A Goldberg	1634				
The MAILING DATE of this communication appe	ars on the cover sheet with the c	orrespondence add	ress			
THE REPLY FILED 03 March 2004 FAILS TO PLACE THIS APPLICATION IN CONDITION FOR ALLOWANCE. Therefore, further action by the applicant is required to avoid abandonment of this application. A proper reply to a final rejection under 37 CFR 1.113 may only be either: (1) a timely filed amendment which places the application in condition for allowance; (2) a timely filed Notice of Appeal (with appeal fee); or (3) a timely filed Request for Continued Examination (RCE) in compliance with 37 CFR 1.114.						
PERIOD FOR RE	EPLY [check either a) or b)]					
a) The period for reply expires 5 months from the mailing date of the final rejection. b) The period for reply expires on: (1) the mailing date of this Advisory Action, or (2) the date set forth in the final rejection, whichever is later. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of the final rejection. ONLY CHECK THIS BOX WHEN THE FIRST REPLY WAS FILED WITHIN TWO MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f). Extensions of time may be obtained under 37 CFR 1.136(a). The date on which the petition under 37 CFR 1.136(a) and the appropriate extension						
fee have been filed is the date for purposes of determining the period of extension and the corresponding amount of the fee. The appropriate extension fee under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the shortened statutory period for reply originally set in the final Office action; or (2) as set forth in (b) above, if checked. Any reply received by the Office later than three months after the mailing date of the final rejection, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
1. A Notice of Appeal was filed on 03 March 2004. Appellant's Brief must be filed within the period set forth in 37 CFR 1.192(a), or any extension thereof (37 CFR 1.191(d)), to avoid dismissal of the appeal.						
2. The proposed amendment(s) will not be entered because:						
(a) ☐ they raise new issues that would require further consideration and/or search (see NOTE below);						
(b) ☐ they raise the issue of new matter (see Note below);						
(c) ☐ they are not deemed to place the application in better form for appeal by materially reducing or simplifying the issues for appeal; and/or						
(d) ☐ they present additional claims without cancel NOTE:	ing a corresponding number of f	inally rejected claim	18.			
3. Applicant's reply has overcome the following reject	tion(s):					
Newly proposed or amended claim(s) would canceling the non-allowable claim(s).		eparate, timely filed	amendment			
5.⊠ The a) affidavit, b) exhibit, or c) request fo application in condition for allowance because: Se	r reconsideration has been cons ee Continuation Sheet.	idered but does NC	T place the			
6. The affidavit or exhibit will NOT be considered becaused by the Examiner in the final rejection.	cause it is not directed SOLELY	to issues which wer	re newly			
7. For purposes of Appeal, the proposed amendmen explanation of how the new or amended claims w	t(s) a)⊡ will not be entered or b rould be rejected is provided belo	o)⊠ will be entered ow or appended.	and an			
The status of the claim(s) is (or will be) as follows:						
Claim(s) allowed: NONE.						
Claim(s) objected to: <u>NONE</u> .						
Claim(s) rejected: <u>1,6-8,11,12 and 15-23</u> .						
Claim(s) withdrawn from consideration: <u>13,14 and 27</u> .						
8. The drawing correction filed on is a) approved or b) disapproved by the Examiner.						
9. Note the attached Information Disclosure Statement(s)(PTO-1449) Paper No(s)						
10.⊠ Other: <u>See Continuation Sheet</u>	Alblig					

U.S. Patent and Trademark Office PTOL-303 (Rev. 11-03) Continuation of 5. does NOT place the application in condition for allowance.

The response argues that the claims are drawn to screening methods of genotypes to determine whether a genotype is associated with increased bipolar affective disorder. The claims as written do not make is clear that the claim is drawn to a screening method. The claim requires using a marker associated with resistance to bipolar disorder. Therefore it is not clear that this is screening for markers associated with resistance to bipolar affective disorder. Applicant argues selection of variants is routine. This is not correct because it is entirely unpredictable if there are variants. In fact, Applicant is attempting to claim this subject matter because it is not routine. It is entirely unpredictable and inventive whether any particular microsatellite marker, translocation, mutation, deletion, splice variant or polymorphism is associated with BPAD. In particular, this unpredictability, combined with the other factors, supports a conclusion of undu experimentation.

Unlike the simple screening assay in Wands itself, where experimental success was assured so long as sufficient resources were expended, since eventually an antibody producing cell would be isolated, here there is no assurance or even likelihood of success, since there is no reason to believe that other polymorphisms necessarily exist which have the desired correlation. At the time of the invention, it is speculative and without evidentiary basis to predict if there will be any results from the screening for additional polymorphisms, unlike Wands where it is not only possible but expected that results will be achieved. Here, there is no expectation that other polymorphisms associated with BPAD will be found. The claimed invention is an idea for future investigation or research. The exact purpose of the method is to determine whether additional genotypes are associated with BPAD.

It is acknowledged that there is at least one marker within each of the claimed regions which is correlated, however there are also severa markers which are not correlated. It is noted that each of these regions is extremely large. For example the region on chromosome 4 is a range of 33.3cM and 42.8cM. These are much larger than the attorney arguments stating that "one would reasonably expect that marker located at about 10cM from a marker would be linked to that marker. It is noted that this is attorney arguments. The submission of the post filing date article to try and establish the current practice in the field is not appropriate (page 10 of response). The enablement of an invention is determined at the time of filing. The instant filing date is 1997. Therefore, the state of the art in 2003 is not at the time the invention was made. As provided by MPEP 2164.05, "To overcome a prima facie case of lack of enablement, applicant must demonstrate by argument and/or evidence that the disclosure, as filed, would have enabled the claimed invention for one skilled in the art at the time of filing."

The response asserts that the fact that markers within a region of interest may not appear significantly correlated with BPAD does not cas doubt on the conclusion that the region is correlated with BPAD. This argument has been thoroughly reviewed, a found persuassive based upon further review of the art and the arguments.

The response correctly argues that the specification, not the claims, provides the general method for screening individuals for particular predispositions. The basic method is well know method for comparing and determining whether an association exists, however, the outcome of each of the markers encompassed by the claims is unpredictable and undue experimentation. Just because the method can be performed does not enable the skilled artisan to make the specific genotypes. The nature of alleles and markers are such that the general knowledge in the art concerning one allele does not provide any indication of how the structure of one alleles is representative of unknown alleles. They are variant structures, and in the present state of the art the structure of one does not provide guidance to the structure of others and the enablement of the others.

With respect to the description issue, the specification fails to provide any deletions, translocations, SNPs, or mutations which are within the scope of the genus. While the response does not appear to focus on the description issue. The specification has not described and was not in possession of a representative number of markers which were linked to a locus associated with resistance to bipolar disorder i the 33cm region of D4S431 and D4S404.

Continuation of 10. Other: Upon review of the formal matters of the application, there is no clear indication of the relationship of the application to 08/827,568 in the first line of the specification or the ADS sheet. Correction is required by 37 CFR 1.78(a)(2) and 1.76. A specific relationship between applications and the status of the applications is required by the provision. It is further noted that Failure to timely submitthe reference required by 37 CFR 1.78(a)(2)(i)is considered a waiver of any benefitclaim under 35 U.S.C.§ 120,,121 or 365(c)unless a petition to accept an unintentionallydelayed claim under 37 CFR 1.78(a)(3),the surcharge set forth in 37 CFR 1.17(t),and therequired reference,including the relationship of the applications (unless previouslysubmitted)are filed. For example, if a benefit claim is submitted without the specific relationship between the nonprovisional applications before the expiration of the period,and the specific relationship between the nonprovisional applications is subsequentlysubmitted after the expiration of the period,a petition and the surcharge would berequired.